



09/463681

01/31/00

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE  
REQUEST FOR FILING NATIONAL PHASE OF  
PCT APPLICATION UNDER 35 U.S.C. 371 AND 37 CFR 1.494 OR 1.495**

To: Asst. Commissioner of Patents  
and Trademarks  
Washington, D.C. 20231

(Our Deposit Account No. 03-3975)

TRANSMITTAL LETTER TO THE UNITED STATES  
DESIGNATED/ELECTED OFFICE (DO/EO/US)

Atty Dkt: PM 266043 /H 33 430  
M# /Client Ref.

From: Pillsbury Madison & Sutro LLP, IP Group:

Date: January 31, 2000

This is a **REQUEST** for FILING a PCT/USA National Phase Application based on:

- |   |  |   |
|---|--|---|
| 1. International Application<br><br><u>PCT/EP98/04614</u> ✓<br>↑ country code | 2. International Filing Date<br><br>23 July 1998 ✓<br>Day MONTH Year | 3. Earliest Priority Date Claimed<br><br>1 August 1997 ✓<br>Day MONTH Year<br>(use item 2 if no earlier priority) |
|---|--|---|

4. Measured from the earliest priority date in item 3, this PCT/USA National Phase Application Request is being filed within:

- (a) ☐ 20 months from above item 3 date      (b) ☒ 30 months from above item 3 date,

(c) Therefore, the due date (unextendable) is February 1, 2000

5. Title of Invention PROCESS FOR THE APPLICATION OF A PROTECTIVE DECORATIVE LAMINAR STRUCTURE

6. Inventor(s) BLUM, et al

Applicant herewith submits the following under 35 U.S.C. 371 to effect filing:

7. ☒ Please immediately start national examination procedures (35 U.S.C. 371 (f)).
8. ☐ A copy of the International Application as filed (35 U.S.C. 371(c)(2)) is transmitted herewith (file if in English) but, if in foreign language, file only if not transmitted to PTO by the International Bureau) including:
- a. ☐ Request;
- b. ☐ Abstract;
- c. ☐ pgs. Spec. and Claims;
- d. ☐ sheet(s) Drawing which are ☐ informal ☐ formal of size ☐ A4 ☐ 11"
9. ☒ A copy of the International Application has been transmitted by the International Bureau.
10. A translation of the International Application into English (35 U.S.C. 371(c)(2))
- a. ☒ is transmitted herewith including: (1) ☐ Request; (2) ☒ Abstract;
- (3) 29 pgs. Spec. and Claims;
- (4)        sheet(s) Drawing which are: ☐ informal ☐ formal of size ☐ A4 ☐ 11"
- b. ☐ is not required, as the application was filed in English.
- c. ☐ is not herewith, but will be filed when required by the forthcoming PTO Missing Requirements Notice per Rule 494(c) if box 4(a) is X'd or Rule 495(c) if box 4(b) is X'd.
- d. ☐ Translation verification attached (not required now).

RE: USA National Filing of PCT/EP98/04614

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11. ☒ **PLEASE AMEND** the specification before its first line by inserting as a separate paragraph:  
 a. ☒ --This application is the national phase of international application PCT/EP98/04614 filed July 23, 1998 which designated the U.S.--  
 b. ☐ --This application also claims the benefit of U.S. Provisional Application No. 60/\_\_\_\_, filed \_\_\_\_--
12. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)), i.e., **before 18th month from first priority date above in item 3, are transmitted herewith (file only if in English) including:**
13. ☒ PCT Article 19 claim amendments (if any) have been transmitted by the International Bureau
14. ☐ Translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)), i.e., of claim amendments made before 18th month, is attached (**required by 20th month from the date in item 3 if box 4(a) above is X'd, or 30th month if box 4(b) is X'd, or else amendments will be considered canceled.**)
15. **A declaration of the inventor** (35 U.S.C. 371(c)(4))  
 a. ☒ is submitted herewith ☒ Original ☐ Facsimile/Copy  
 b. ☐ is not herewith, but will be filed when required by the forthcoming PTO Missing Requirements Notice per Rule 494(c) if box 4(a) is X'd or Rule 495(c) if box 4(b) is X'd.
16. **An International Search Report (ISR):**  
 a. Was prepared by ☐ European Patent Office ☐ Japanese Patent Office ☐ Other  
 b. ☒ has been transmitted by the international Bureau to PTO.  
 c. ☐ copy herewith (\_\_\_\_ pg(s).) ☐ plus Annex of family members (\_\_\_\_ pg(s)).
17. **International Preliminary Examination Report (IPER):**  
 a. ☒ has been transmitted (if this letter is filed after 28 months from date in item 3) in English by the International Bureau with Annexes (if any) in original language.  
 b. ☐ copy herewith in English.  
 c. 1 ☐ IPER Annex(es) in original language ("Annexes" are amendments made to claims/spec/drawings during Examination) including attached amended:  
 Specification/claim pages #\_\_\_\_ claims #\_\_\_\_  
 Dwg Sheets #\_\_\_\_  
 c. 2 ☐  
 d. ☐ Translation of Annex(es) to IPER (**required by 30th month due date, or else annexed amendments will be considered canceled.**)
18. **Information Disclosure Statement** including:  
 a. ☐ Attached Form PTO-1449 listing documents  
 b. ☐ Attached copies of documents listed on Form PTO-1449  
 c. ☒ A concise explanation of relevance of ISR references is given in the ISR.
19. ☐ **Assignment** document and Cover Sheet for recording are attached. Please mail the recorded assignment document back to the person whose signature, name and address appear at the end of this letter.
20. ☐ Copy of Power to IA agent.
21. ☐ **Drawings** (complete only if 8d or 10a(4) not completed): \_\_\_\_ sheet(s) per set: ☐ 1 set informal; ☐ Formal of size ☐ A4 ☐ 11"
22. ☐ \_\_\_\_ (No.) **Verified Statement(s)** establishing "small entity" status under Rules 9 & 27
23. **Priority** is hereby claimed under 35 U.S.C. 119/365 based on the priority claim and the certified copy, both filed in the International Application during the international stage based on the filing in (country) GERMANY of:  

	<u>Application No.</u>	<u>Filing Date</u>		<u>Application No.</u>	<u>Filing Date</u>
(1)	<u>197 33 312.5</u>	<u>August 1, 1997</u>	(2)	_____	_____
(3)	_____	_____	(4)	_____	_____
(5)	_____	_____	(6)	_____	_____

 a. ☒ See Form PCT/IB/304 sent to US/DO with copy of priority documents. If copy has not been received, **please proceed promptly to obtain same from the IB.**

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24. Attached:

25. Preliminary Amendment: attached

25.5 Per Item 17.c2, **cancel original** pages #\_\_, claims #\_\_, Drawing Sheets #26. **Calculation of the U.S. National Fee (35 U.S.C. 371 (c)(1)) and other fees is as follows:**Based on amended claim(s) per above item(s) ☐ 12, ☐ 14, ☐ 17, ☒ 25, ☐ 25.5 (hilitte)

Total Effective Claims	10	minus 20 =	0	x \$18/\$9	=	\$0	966/967
Independent Claims	1	minus 3 =	0	x \$78/\$39	=	\$0	964/965
If any proper (ignore improper) Multiple Dependent claim is present,				add \$260/\$130	+	0	968/969

BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(4)): →→ BASIC FEE REQUIRED, **NOW** →→→→A. If country code letters in item 1 are **not** "US", "BR", "BB", "TT", "MX", "IL", "NZ", "IN" or "ZA" ↓

See item 16 re: ↓

1. Search Report was <b>not</b> prepared by EPO or JPO -----	add \$970/\$485	960/961
2. Search Report was prepared by EPO or JPO -----	add \$840/\$420	970/971
	<b>+840</b>	

SKIP B, C, D AND E UNLESS country code letters in item 1 are "US", "BR", "BB", "TT", "MX", "IL", "NZ", "IN" or "ZA" ↓

→ <input type="checkbox"/> B. If USPTO did not issue <b>both</b> International Search Report (ISR) <b>and</b> (if box 4(b) above is X'd) the International Examination Report (IPER), -----	add \$970/\$485	+0	960/961
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→ <input type="checkbox"/> C. If USPTO issued ISR but not IPER (or box 4(a) above is X'd), -----	add \$690/\$345	+0	958/959
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→ <input type="checkbox"/> D. If USPTO issued IPER but IPER Sec. V boxes <b>not all</b> 3 YES, -----	add \$670/\$335	+0	956/957
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→ <input type="checkbox"/> E. If international preliminary examination fee was paid to USPTO and Rules 492(a)(4) and 496(b) <b>satisfied</b> (IPER Sec. V <b>all</b> 3 boxes YES for <b>all</b> claims), -----	add \$96/\$48	+0	962/963
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27. **SUBTOTAL = \$840**

28. If Assignment box 19 above is X'd, add Assignment Recording fee of ---\$40 +0 (581)

29. Attached is a check to cover the ----- **TOTAL FEES \$840**

Our Deposit Account No. 03-3975

Our Order No. 71461

266043

C#

M#

**CHARGE STATEMENT:** The Commissioner is hereby authorized to charge any fee specifically authorized hereafter, or any missing or insufficient fee(s) filed, or asserted to be filed, or which should have been filed herewith or concerning any paper filed hereafter, and which may be required under Rules 16-18 and 452 (missing or insufficient fee only) now or hereafter relative to this application and the resulting Official document under Rule 20, or credit any overpayment, to our Account/Order Nos. shown above for which purpose a duplicate copy of this sheet is attached.

This CHARGE STATEMENT **does not authorize** charge of the issue fee until/unless an issue fee transmittal form is filedPillsbury Madison & Sutro LLP  
Intellectual Property Group1100 New York Avenue, NW  
Ninth Floor  
Washington, DC 20005-3918  
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By Atty: Ruth N. Morduch

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## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

BLUM et al.

Group Art Unit: unknown

Appln. No. 09/unassigned

Examiner: unassigned

Filed: Herewith

For: PROCESS FOR THE APPLICATION OF A PROTECTIVE  
AND DECORATIVE LAMINAR STRUCTURE

\* \* \* \* \*

January 31, 2000

## PRELIMINARY AMENDMENT

Hon. Commissioner of Patents  
and Trademarks  
Washington, D.C. 20231

Sir:

Prior to examination, please amend the above-identified application as  
follows.

## IN THE SPECIFICATION:

Page 1, line 3, insert the following centered heading:

**--FIELD OF THE INVENTION--;**

line 8, insert the following centered heading:

**--BACKGROUND OF THE INVENTION--;** and

line 35, insert the following centered heading:

**--SUMMARY OF THE INVENTION--.**

Page 4, line 8, insert the following centered heading:

**--DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS--.**Page 28, line 1, change "Patent Claims" to **--WHAT IS CLAIMED IS--.**

Page 30, line 1, change "Abstract" to the following centered heading:

**--ABSTRACT OF THE DISCLOSURE--.**

**IN THE CLAIMS:**

Please amend the claims as follows:

Claim 1, line 3, change "characterised in that" to --without using spray application, wherein--.

line 8, delete "without spraying".

Claim 2, line 1, change "characterised in that" to --wherein--.

Claim 3, lines 1 and 2, change "or 2, characterised in that" to --wherein--.

Claim 4, lines 1 and 2, change "any one of claims 1, 2 or 3, characterised in that" to --claim 1, wherein--.

Claim 5, lines 1 and 2, change "any one of claims 1 to 4, characterised in that" to --claim 1, wherein--.

Claim 6, lines 1 and 2, change "any one of claims 1 to 5, characterised in that" to --claim 1, wherein--.

Claim 7, lines 1 and 2, change "any one of claims 1 to 5, characterised in that" to --claim 1, wherein--.

**BLUM et al. - New U.S. Patent Appln.**

Claim 8, lines 1 and 2, change "any one of claims 1 to 7, characterised in that it is used for" to --claim 1, comprising--.

Claim 9, line 2, change "any one of claims 1 to 8" to --claim 1--.

Claim 10, line 3, change "any one of claims 1 to 8" to --claim 1--.

**REMARKS**

The amendments to the specification were made to add the subheadings required by Patent Office practice.

The amendments to the claims were made to eliminate the multiple dependencies. No new matter was intended to be added, nor is any new matter believed to have been added. Accordingly, an early action on the merits is earnestly solicited.

Respectfully submitted,

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Process for the application of a protective and decorative  
laminar structure

5 This invention relates to a process for providing the surface of electrically conductive three-dimensional substrates with a protective and decorative laminar structure and to the resultant laminar structures.

10 Modern automotive original lacquer coatings usually consist of a base coat/clear topcoat lacquer coating, which is applied onto a body which has been electrophoretically primed and coated with surfacer. In such coatings, the surfacer, base coat lacquer and clear lacquer are applied by spraying. Large quantities of problematic lacquer sludge  
15 are formed during the spray application in the lacquer coating process. The lacquer coating process demands elaborate logistics on the part of the vehicle manufacturer due not only to the multiple lacquer layers of different lacquer materials to be applied but also to the numerous  
20 shades in which some of the lacquer layers are produced. The base lacquer layers are thus applied in various different shades, as are the surfacer layers too by many automotive manufacturers.

25 In the Federal Republic of Germany, for example, vehicles marked as taxis are of a prescribed shade. Vehicles which have been used as taxis are difficult to sell as they can be recognised as former taxis. For this reason, conventional cars not painted in the taxi shade have been provided  
30 with a bonded plastic film of the taxi shade which hides the actual shade of the multi-layer lacquer coating. This film may subsequently be removed once the taxi has completed its service, so revealing the actual lacquer coating again.

35 The object of the invention is to provide a logistically straightforward process which avoids lacquer waste and permits the surface of three-dimensional substrates, in particular motor vehicles, to be provided with a protective

coating and decorative finish. In particular, it should be possible to perform the process in an environmentally friendly and efficient manner.

- 5 It has been found that this object may be achieved by a process for the production of a three-dimensional substrate provided with a protective and decorative laminar structure, which process is characterised in that a primer layer of a coating composition (I) which is electrically  
10 conductive in the stoved state is applied without spraying onto an electrically conductive substrate and stoved, whereupon a substrate not yet in the desired three-dimensional shape is shaped, whereupon a second coating layer of an electrophoretically depositable coating  
15 composition (II) is electrophoretically deposited and stoved and whereupon a plastic film is applied.

- In the process according to the invention, the conductive primer layer is in particular applied by brushing, roller  
20 application, dipping or flow coating. The conductive primer layer is particularly preferably applied using the coil coating process, by autophoretic deposition or electrophoretic deposition. The conductive primer layer may be applied onto the entire surface, i.e. both sides, of the  
25 substrate. The substrate may be in the desired three-dimensional shape even before application of the primer layer. It is also possible not to shape the substrate three-dimensionally until the conductive primer layer has been applied and stoved, wherein it has optionally  
30 previously been stamped or cut. Shaping may be achieved by the usual processes familiar to the person skilled in the art for the particular substrate, for example by deep drawing. The substrate may also be in the form of individual components which may be combined into an  
35 assembly before application of the second coating layer, wherein the individual components may already have been provided with the primer layer.



Various coating compositions (I) may be used to apply the coating layer which is electrically conductive in the stoved state, wherein no spray application methods are used according to the invention.

5

A first preferred embodiment of the present invention comprises a process for the production of a protective and decorative laminar structure on a three-dimensional, electrically conductive substrate, in which a primer layer  
 10 of a primer coating composition (Ia) which is electrically conductive in the stoved state is applied onto both sides of a sheet metal coil using the coil coating process and stoved, sheet metal components are subsequently stamped out from the coil and shaped by deep drawing and optionally  
 15 joined together into an assembly, whereupon a second coating layer of an electrophoretically depositable coating composition (II) is electrophoretically deposited and stoved and whereupon a plastic film is subsequently applied.

20

A second preferred embodiment of the present invention comprises a process for the production of a protective and decorative laminar structure on a three-dimensional, electrically conductive substrate having an  
 25 autophoretically coatable surface, in which a primer layer of an electrophoretically depositable coating composition (Ib) which is electrically conductive in the stoved state is autophoretically deposited and stoved, whereupon a further coating layer of an electrophoretically depositable  
 30 coating composition (II) is electrophoretically deposited and stoved and whereupon a plastic film is subsequently applied.

A third and particularly preferred embodiment of the  
 35 present invention comprises a process for the production of a protective and decorative laminar structure on a three-dimensional, electrically conductive substrate, in which a

primer layer of an electrophoretically depositable coating composition (Ic) which is electrically conductive in the stoved state is electrophoretically deposited and stoved, whereupon a further coating layer of an electrophoretically depositable coating composition (II) other than (Ic) is electrophoretically deposited and stoved and whereupon a plastic film is subsequently applied.

In the first preferred embodiment of the process according to the invention, a primer layer of a primer coating composition (Ia) is produced of such a film thickness that an electrically conductive coating layer is obtained in the stoved state. After stamping and shaping of the sheet metal components primed in this manner and optionally joined together into an assembly, in particular an automotive body, the second layer may then be applied by electrophoretic deposition from an electrophoretically depositable aqueous coating composition (II).

The primer coating compositions (Ia) in particular comprise liquid coating compositions which may be applied by the coil coating process. The coating compositions may be aqueous or based on organic solvents. They may be physically drying. They are generally crosslinkable by the formation of covalent bonds. The coating compositions may be self-crosslinking or extrinsically crosslinking systems.

The primer coating compositions (Ia) usable for the production of the primer layer in the process according to the invention contain one or more film-forming binders. They may optionally also contain crosslinking agents, in particular in the event that the binders are not self-crosslinking or are physically drying or thermoplastic. No particular restrictions apply to either the binder component or the crosslinking agent component; resins conventional in lacquers and familiar to the person skilled in the art may be used. Polyester, polyurethane, epoxy

and/or polymer resins may, for example, be used as the film-forming binders.

Selection of the optionally present crosslinking agent is determined by the functionality of the binders, i.e. the crosslinking agents are selected from among conventional crosslinking agents familiar to the person skilled in the art in such a manner that they have a reactive functionality complementary to the functionality of the binder. Examples of such complementary functionalities between binders and crosslinking agents are: hydroxyl/methylol ether, hydroxyl/blocked isocyanate. Providing that they are mutually compatible, two or more such complementary functionalities may simultaneously be present. The crosslinking agents optionally used in the primer coating compositions (Ia) may be present individually or as a mixture.

It is preferred in the process according to the invention for the primer coating compositions (Ia) applied in the coil coating process to contain constituents which confer electrical conductivity and impart to the primer layer in the stoved state a sufficiently low specific resistance, generally of below  $10^8 \Omega \text{ cm}$ , for example between  $10^3$  and  $10^8 \Omega \text{ cm}$ , for the electrophoretic deposition of a further coating layer of an electrophoretically depositable coating composition (II). This also ensures that the primer coating layer is spot-weldable. Examples of such components which confer electrical conductivity are particulate electrical conductors or semi-conductors, as are conventional for this purpose and familiar to the person skilled in the art, for example iron oxide black, graphite, conductive carbon black, metal powders, for example of aluminium, zinc, copper or stainless steel, molybdenum disulfide, special conductive pigments based on mica flakes, for example mica flakes coated with antimony-doped tin dioxide. The constituents which impart electrical conductivity are

present in the primer coating composition (Ia) in a quantity such that the desired specific resistance of the primer coating layer in the stoved state is achieved. Relative to the solids content of the primer coating composition (Ia), the proportion of the constituent or constituents which impart(s) electrical conductivity is, for example, between 1 and 30 wt.%. The proportion may readily be determined by the person skilled in the art; it is dependent, for example, on the specific weight, specific electrical conductivity and particle size of the constituents used to impart electrical conductivity. One or more of these constituents may be present in combination.

The primer coating compositions (Ia) may contain pigments and/or extenders. Pigments or extenders which may be considered are, for example, conventional inorganic or organic pigments and/or extenders. Examples are carbon black, titanium dioxide, iron oxide pigments, kaolin, talcum, silicon dioxide and in particular anti-corrosion pigments, such as zinc chromate, strontium chromate, lead silicate, zinc phosphate, aluminium phosphate, strontium aluminium polyphosphate hydrate.

The primer coating compositions (Ia) may furthermore contain conventional additives. Examples of these are conventional lacquer additives, such as wetting agents, dispersion auxiliaries, emulsifiers, levelling agents, corrosion inhibitors, anti-foaming agents and in particular lubricants, for example waxes such as polyethylene wax, molybdenum disulfide, graphite.

Primer coating compositions (Ia) suitable for the production of primer layers, applicable using the coil coating process and containing no special or adequate quantities of constituents which impart electrical conductivity, are those, from which primer layers may be produced while retaining a low dry film layer thickness of

for example 1 to 3  $\mu\text{m}$ , which exhibit sufficient electrical conductivity in the stoved state for deposition of a subsequent coating layer of an electrophoretically depositable coating composition (II) and, preferably, also spot-weldability.

The primer coating compositions (Ia) preferably usable for the production of primer layers, applicable using the coil coating process and which may be applied in greater dry film thicknesses of for example 2 to 15  $\mu\text{m}$ , preferably of 3 to 10  $\mu\text{m}$  by virtue of their content of constituents conferring electrical conductivity, are those coating compositions from which primer layers may be produced which exhibit a sufficiently low specific electrical resistance for the deposition of a subsequent coating layer of an electrophoretically depositable coating composition (II) and which are preferably spot-weldable.

Substrates used for the first embodiment of the process according to the invention are metal sheets wound into coils, for example having a sheet thickness typical for bodywork construction of 0.7 to 1.3 mm, for example made from iron, zinc, aluminium or corresponding alloys. Galvanised sheet metal, for example galvanised sheet steel is preferred. The metal surfaces may be pretreated, for example by phosphating, and optionally be provided with passivation.

In the coil coating process, the primer coating composition (Ia) is rolled onto both sides of the sheet metal substrate. If the primer coating composition (Ia) is one which contains no special constituents which confer electrical conductivity, it is applied in a sufficiently low dry film thickness to ensure sufficient electrical conductivity for electrophoretic deposition of a further coating layer of an electrophoretically depositable coating composition (II), i.e. of a dry film thickness which is not

yet electrically insulating, for example in a dry film thickness of 1 to 3  $\mu\text{m}$ . If, in contrast, it is a primer coating composition (Ia) preferred in the process according to the invention which contains constituents conferring electrical conductivity which impart a sufficiently low specific resistance to the first coating layer in the stoved state for electrophoretic deposition of a further coating layer of an electrophoretically depositable coating composition (II), the primer coating composition (Ia) is applied in a dry film thickness of, for example, 2 to 15  $\mu\text{m}$ , preferably of 3 to 10  $\mu\text{m}$ . Stoving of the primer coating layer proceeds rapidly, for example within 30 to 60 seconds, at elevated temperature, for example at 300 to 370°C, wherein maximum object temperatures (PMT, peak metal temperatures) of, for example, between 200 and 260°C are achieved, which stoving may proceed, for example, in a convection oven.

Sheet metal components of the desired shape are conventionally stamped and then shaped, for example by deep drawing, from the metal coil provided on both sides with the stoved primer layer. The shaped components may then, optionally with components made from other materials, be joined together into an assembly, in particular a vehicle body, for example by adhesive bonding and/or clinching and/or bolting and/or preferably welding, for example spot welding. The components made from other materials may, for example, be differently coated or uncoated metals or plastics, preferred components being those having a conductive and electrophoretically coatable surface.

In the second preferred embodiment of the process according to the invention, a primer layer of an autophoretically depositable coating composition (Ib) is autophoretically deposited and stoved, wherein in the stoved state an electrically conductive coating layer is obtained. The second layer may then be applied by electrophoretic

deposition from an electrophoretically depositable aqueous coating composition (II).

- Autophoretically depositable coating compositions (Ib)
- 5 which are known *per se* may be used as the autophoretically coating compositions (Ib), wherein these compositions contain constituents which impart to the primer layer in the stoved state a sufficiently low specific resistance for electrophoretic deposition of a further coating composition
- 10 of an electrophoretically depositable coating composition (II).

- The autophoretically depositable coating compositions (Ib) are coating compositions based on aqueous binder
- 15 dispersions having a negative surface charge of the binder particles. By virtue of their generally acidic pH value of for example between 1 and 6, preferably between 1.5 and 5.0, and their generally oxidising nature, autophoretic lacquers are capable of attacking sufficiently non-noble
- 20 metal surfaces, so liberating the corresponding metal ions. If this results in the formation of a concentration of metal ions in the vicinity of the metal surface which is sufficient to destabilise and coagulate the binder particles dispersed in the aqueous phase, then a coating
- 25 film is deposited on the metal surface.

- The autophoretically depositable coating compositions (Ib) generally have a low solids content of, for example, up to 20 wt.%, wherein the lower limit is generally, for example,
- 30 5 wt.%, and the upper limit for example 10 wt.%. In addition to the autophoretically depositable film-forming binder, the compositions contain water, acid and electrically conductive constituents, as well as in general oxidising agents, optionally together with crosslinking
- 35 agents for the binder, extenders, pigments and conventional lacquer additives.

The autophoretically depositable coating compositions (Ib) may be physically drying or crosslinkable with the formation of covalent bonds. The autophoretic lacquers (Ib) which crosslink with the formation of covalent bonds may be self-crosslinking or extrinsically crosslinking systems.

The autophoretic lacquers (Ib) usable for the production of the primer layer in the process according to the invention contain one or more film-forming binders containing *per se* neutral or anionic groups. In particular in the event that the binders are not self-crosslinking or are physically drying or thermoplastic, the lacquers may optionally also contain crosslinking agents. Binders and optionally present crosslinking agents are in the form of an aqueous dispersion having a negative surface charge of the dispersed particles. The negative surface charge stabilises the dispersed particles in the aqueous phase. The negative surface charge may, for example, originate from anionic groups in the binder and/or in particular in the case of *per se* neutral binders, from an anionic emulsifier for the binder and the crosslinking agents. Examples of anionic groups in the binder are anionic groups of the binder *per se*, for example carboxyl groups or sulfonate groups, and/or anionic residues from production of the binder, for example from the production of a *per se* neutral binder. Examples of anionic residues from production of the binder are sulfate groups as residues remaining in the binder from a free-radical polymerisation initiated by peroxydisulfate. No restrictions apply to either the binder component or to the crosslinking agent component; resins conventional in lacquers and familiar to the person skilled in the art may be used. Polyester, polyurethane, epoxy and/or polymer resins may, for example, be used as the film-forming binder. Polymer resins, i.e. binders produced by free-radical polymerisation, in particular by emulsion polymerisation or seed polymerisation, are particularly preferred. Conventional aqueous, thermoplastic polymer



dispersions (lattices) which contain homo- or copolymers of olefinically unsaturated monomers having glass transition temperatures of for example between 0 and 100°C as the disperse phase are particularly preferred. Examples of

5 suitable olefinically unsaturated monomers for the synthesis of such homo- and copolymers are (meth)acrylic acid esters, such as for example methyl (meth)acrylate, ethyl (meth)acrylate, n-butyl (meth)acrylate, hydroxyethyl (meth)acrylate, hydroxypropyl (meth)acrylate, ethylhexyl

10 (meth)acrylate; glycidyl esters of unsaturated carboxylic acids, such as for example glycidyl (meth)acrylate; (meth)acrylamide, (meth)acrylonitrile, monomers containing anionic groups, such as alkali metal, for example sodium, 2-sulfoethylmethacrylate, (meth)acrylic acid; as well as

15 monomers having no further functional groups or other functional groups, such as for example ethylene, styrene, vinyl chloride, vinylidene chloride, vinyl acetate, butadiene. Examples of aqueous polymer dispersions suitable for the purposes of the invention are styrene/butadiene,

20 butadiene/acrylonitrile, vinyl chloride, ethylene/vinyl acetate and vinylidene chloride copolymer dispersions.

(Meth)acrylic here denotes acrylic and/or methacrylic.

25 Selection of the optionally present crosslinking agent is determined by the functionality of the binders, i.e. the crosslinking agents are selected from among conventional crosslinking agents familiar to the person skilled in the art in such a manner that they have a reactive

30 functionality complementary to the functionality of the binders. Examples of such complementary functionalities between binders and crosslinking agents are: hydroxyl/methylol ether, hydroxyl/blocked isocyanate. Providing that they are mutually compatible, two or more

35 such complementary functionalities may simultaneously be present in an autophoretic lacquer (Ib). The crosslinking

agents optionally used in the autophoretic lacquers (Ib) may be present individually or as a mixture.

- The autophoretically depositable coating compositions (Ib)
- 5 contain one or more free acids in a quantity to establish the above-stated pH range and preferably also one or more oxidising agents. Examples of preferred acids are inorganic acids, such as hydrochloric acid, phosphoric acid, sulfuric acid, nitric acid, and organic acids, such as formic acid,
- 10 acetic acid. Hydrofluoric acid is particularly preferred. Examples of preferred oxidising agents are chromate, dichromate, bromate, chlorate, perchlorate, permanganate, persulfate, peroxydisulfate. Hydrogen peroxide is particularly preferred.
- 15 A particularly preferred combination of acid and oxidising agent is the combination hydrofluoric acid/hydrogen peroxide.
- 20 The autophoretically depositable coating compositions (Ib) contain constituents which confer electrical conductivity. They are intended to impart to the primer layer in the stoved state a sufficiently low specific resistance, generally of below  $10^8 \Omega \text{ cm}$ , for example between  $10^3$  and
- 25  $10^8 \Omega \text{ cm}$ , for the electrophoretic deposition of a further coating layer of an electrophoretically depositable coating composition (II). Examples of such components are particulate electrical conductors or semi-conductors, as are conventional for this purpose and familiar to the
- 30 person skilled in the art, for example iron oxide black, graphite, conductive carbon black, metal powders, for example of copper or stainless steel, molybdenum disulfide, special conductive pigments based on mica flakes, for example mica flakes coated with antimony-doped tin dioxide.
- 35 The constituents which impart electrical conductivity are present in the autophoretically depositable coating composition (Ib) in a quantity such that the desired

specific resistance of the coating layer deposited therefrom is achieved in the stoved state. Relative to the solids content of the autophoretically deposited coating composition (Ib), the proportion of the constituent or constituents which impart(s) electrical conductivity is, for example, between 1 and 30 wt.%. The proportion may readily be determined by the person skilled in the art; it is dependent, for example, on the specific weight, specific electrical conductivity and particle size of the constituents used to impart electrical conductivity. One or more of these constituents may be present in combination.

The autophoretically depositable coating compositions (Ib) may contain pigments and/or extenders. Pigments which may be considered are, for example, conventional inorganic or organic pigments, in particular acid-resistant pigments and/or extenders. Examples are carbon black, titanium dioxide, iron oxide pigments, kaolin, talcum or silicon dioxide as well as anti-corrosion pigments, Pigments, extenders and constituents conferring electrical conductivity to the stoved autophoretic lacquer (Ib) may, for example, be used in formulating the autophoretic lacquer (Ib) as pigment suspensions (pigment slurries) or ground pigment preparations in water and/or organic, water-miscible solvents, such as glycols, for example ethylene glycol, propylene glycol; alcohols, for example sec.-butanol and hexanol; glycol ethers, for example ethoxypropanol, methoxypropanol and butoxyethanol. Such ground pigment preparations are commercially available and are offered for sale, for example, by the company Hoechst under the name Colanyl®.

The autophoretically depositable coating compositions (Ib) usable in the process according to the invention may furthermore contain conventional additives. Examples of these are conventional lacquer additives, such as wetting

- agents, anionic and/or nonionic emulsifiers, protective colloids, levelling agents, corrosion inhibitors, plasticisers, anti-foaming agents, solvents for example as film-forming auxiliaries, light stabilisers, fluorides, in particular for example iron trifluoride, hydrogen fluorides, complex fluorine anions, for example tetrafluoroborates, hexafluorozirconates, hexafluorotitanates.
- 10 Substrates for the second preferred embodiment of the process according to the invention are electrically conductive, autophoretically coatable metal surfaces, in particular metal assemblies, preferably unfinished vehicle bodies having metallic surfaces. These may have been
- 15 manufactured from a single metal or constructed using composite methods from two or more metallic materials and/or from plastic components provided with a suitable metal layer. Metallic surfaces which may be considered are conventional metal surfaces familiar to the person skilled
- 20 in the art, for example of iron, zinc, aluminium or corresponding alloys, as well as, for example, galvanised steel surfaces, which may be attacked by autophoretic lacquer systems, in particular at an acidic pH of for example between 1 and 6 with the liberation of metal ions.
- 25 The metal surfaces may be pretreated, for example by phosphating, and optionally be provided with passivation. It should be noted in this connection that the phrase "unfinished vehicle bodies" in particular includes not only unfinished motor vehicle bodies but also, for example, the
- 30 components thereof and motor vehicle chassis having visible surfaces.
- The primer layer of an autophoretically depositable coating composition (Ib) is deposited autophoretically in the
- 35 conventional manner onto these substrates, preferably using the dipping process, preferably in a dry film thickness of, for example, 5 to 25  $\mu\text{m}$ , particularly preferably of 10 to

20  $\mu\text{m}$ . Before subsequent coating with the electrophoretically depositable coating composition (II), the autophoretically deposited coating layer may be post-treated with water and/or optionally with special solutions, for example rinsed, before stoving. (In connection with the application of the autophoretically depositable coating composition, depending upon the nature of the autophoretic lacquer system, the term "stoving" may mean stoving with chemical crosslinking of the autophoretically deposited coating layer or a purely physical drying at elevated temperature, for example with fusion or sintering and formation of a continuous autophoretic lacquer coating). Stoving proceeds at the temperatures adapted to the autophoretic lacquer system used of, for example, between 80 and 190°C, preferably between 100 and 160°C.

In the third, particularly preferred embodiment of the process according to the invention, a primer layer of an electrophoretically depositable coating composition (Ic) is electrophoretically deposited and stoved, wherein an electrically conductive coating layer is obtained in the stoved state. The second layer may then be applied by electrophoretic deposition of an electrophoretically depositable aqueous coating composition (II) other than (Ic).

Electrophoretically depositable coating compositions (Ic) which may be used are *per se* known anodically or cathodically depositable electrocoating lacquers, wherein it is the case that the electrophoretically depositable coating composition (Ic) contains constituents which impart a sufficiently low specific resistance to the primer layer in the stoved state for the electrophoretic deposition of a further electrophoretically depositable coating composition (II) other than (Ic).

The electrophoretically depositable coating compositions (Ic) are aqueous coating compositions having a solids content of up to 50 wt.%, for example of up to 20 wt.%, wherein the lower limit is, for example, around 10 wt.%.

- 5 The solids content is formed from binders conventional for electrocoating, wherein at least a proportion of the binder bears ionic substituents and/or substituents convertible into ionic groups, optionally together with groups capable of chemical crosslinking, together with optionally present
- 10 crosslinking agents, electrically conductive constituents, extenders, pigments and conventional lacquer additives.

- The ionic groups or groups convertible into ionic groups of the binders may be anionic groups or groups convertible
- 15 into anionic groups, for example acid groups, such as -COOH, -SO<sub>3</sub>H and/or -PO<sub>3</sub>H<sub>2</sub> and the corresponding anionic groups neutralised with bases. They may also be cationic groups or groups convertible into cationic groups, for example basic groups, preferably basic groups containing
  - 20 nitrogen; these groups may be in quaternised form or they are converted into ionic groups with a conventional neutralising agent, for example an organic monocarboxylic acid, such as for example formic acid or acetic acid. Examples are amino, ammonium, for example quaternary
  - 25 ammonium, phosphonium and/or sulfonium groups.

- Anodically depositable electrocoating binders and lacquers containing the conventional anionic groups may, for example, be used. These are, for example, binders based on
- 30 polyesters, epoxy resin esters, (meth)acrylic copolymer resins, maleate oils or polybutadiene oils having a weight average molecular weight (Mw) of, for example, 300-10000 and an acid value of, for example, 35-300 mg of KOH/g. The binders bear, for example, -COOH, -SO<sub>3</sub>H and/or -PO<sub>3</sub>H<sub>2</sub>
  - 35 groups. Once at least a proportion of the acid groups has been neutralised, the resins may be converted into the aqueous phase. The binders may be self-crosslinking or

extrinsically crosslinking. The lacquers may thus also contain conventional crosslinking agents, for example triazine resins, crosslinking agents which contain transesterifiable groups or blocked polyisocyanates.

5

Conventional cathodic electrocoating lacquers based on cationic or basic binders are also usable for the production of the primer layer. Such basic resins are, for example, resins containing primary, secondary and/or

10 tertiary amino groups, the amine values of which are, for example, from 20 to 250 mg of KOH/g. The weight average molecular weight (Mw) of the base resins is preferably 300 to 10000. Examples of such base resins are

15 amino(meth)acrylate resins, aminoepoxy resins, aminoepoxy resins having terminal double bonds, aminoepoxy resins having primary OH groups, aminopolyurethane resins, polybutadiene resins containing amino groups or modified epoxy resin/carbon dioxide/amine reaction products. These

20 base resins may be self-crosslinking or they are used mixed with known crosslinking agents. Examples of such crosslinking agents are amino resins, blocked polyisocyanates, crosslinking agents having terminal double bonds, polyepoxy compounds or crosslinking agents which contain transesterifiable groups.

25

The electrocoating lacquer composition (Ic) contains constituents which confer electrical conductivity. They are intended to impart to the primer layer in the stoved state a sufficiently low specific resistance, generally of below

30  $10^8 \Omega \text{ cm}$ , for example between  $10^3$  and  $10^8 \Omega \text{ cm}$ , for the electrophoretic deposition of a further coating layer of the electrophoretically depositable coating composition (II). Examples of such components are particulate

35 electrical conductors or semi-conductors, such as for example iron oxide black, graphite, conductive carbon black, metal powders, for example of aluminium, copper or stainless steel, molybdenum disulfide, special conductive

pigments based on mica flakes, for example mica flakes coated with antimony-doped tin dioxide, or also polymers exhibiting electrical conductivity, such as for example preferably polyaniline. The constituents which impart electrical conductivity are present in the electrocoating lacquer composition (Ic) in a quantity such that the desired specific resistance of the coating layer deposited therefrom is achieved in the stoved state. Relative to the solids content of the autophoretically deposited coating composition (Ic), the proportion of the constituent or constituents which impart(s) electrical conductivity is, for example, between 1 and 30 wt.%. The proportion may readily be determined by the person skilled in the art; it is dependent, for example, on the specific weight, specific electrical conductivity and particle size of the constituents used to impart electrical conductivity. One or more of these constituents may be present in combination.

In addition to the base resins and optionally present crosslinking agent and the constituents present in the electrocoating lacquer composition (Ic) which impart electrical conductivity to the primer layer in the stoved state, the electrocoating lacquer composition (Ic) may contain pigments, extenders and/or conventional lacquer additives. Pigments which may be considered are, for example, conventional inorganic and/or organic colouring pigments, such as for example titanium dioxide, iron oxide pigments, carbon black, phthalocyanine pigments, quinacridone pigments. Examples of extenders are kaolin, talcum or silicon dioxide.

The pigments may be dispersed to yield pigment pastes, for example using known paste resins.

Possible additives are the conventional additives as are in particular known for electrocoating lacquer compositions. Examples of these are wetting agents, neutralising agents,



levelling agents, catalysts, corrosion inhibitors, anti-foaming agents, solvents.

It is preferred in the process according to the invention  
5 to use a cathodic electrocoating lacquer composition as the electrocoating lacquer composition (Ic).

Suitable substrates for the third, particularly preferred  
embodiment of the process according to the invention are  
10 electrically conductive, electrophoretically coatable assemblies, for example metal assemblies. Automotive bodies or components thereof are in particular suitable; they may consist of pretreated or unpretreated metal or plastic which is electrically conductive or provided with an  
15 electrically conductive layer.

The first coating layer of the electrophoretically depositable coating composition (Ic) is deposited  
electrophoretically in the conventional manner onto these  
20 substrates in a dry film thickness of, for example, 5 to 15  $\mu\text{m}$  and stoved at temperatures of, for example, between 130 and 180°C.

The stoved primer layer obtained in the first preferred  
25 embodiment of the process according to the invention using coating composition (Ia) or in the second preferred embodiment of the process according to the invention using coating composition (Ib) or in the third, particularly preferred embodiment of the process according to the  
30 invention using coating composition (Ic) covers the entire surface of the substrate.

The resultant substrate, preferably an automotive body, provided with an electrically conductive, stoved primer  
35 layer having a specific resistance of in particular below  $10^8$ , for example of between  $10^3$  and  $10^8 \Omega \text{ cm}$  is provided with the second coating layer by electrocoating of an

electrophoretically depositable coating composition (II) other than the electrophoretically depositable coating composition (Ic) and stoved. The plastic film may subsequently be applied according to the invention.

5

The second coating layer may or may not influence the decorative impression of a laminar structure produced according to the invention, the second coating layer preferably does have an influence on the decorative effect.

- 10 In this case, the electrophoretically depositable coating compositions (II) are in particular colouring and/or effect coating compositions.

- 15 In the process according to the invention, the second coating layer may be produced using *per se* known anodically or cathodically depositable electrocoating lacquers as the electrocoating lacquer compositions (II) other than the electrocoating lacquer compositions (Ic).

- 20 The electrocoating lacquer compositions (II) are in particular aqueous coating compositions having a solids content of up to 50 wt.%, for example of up to 20 wt.%, wherein the lower limit is, for example, around 10 wt.%. The solids content is formed from binders conventional for  
25 electrocoating, wherein at least a proportion of the binder bears ionic substituents and/or substituents convertible into ionic groups, optionally together with groups capable of chemical crosslinking, together with optionally present crosslinking agents, extenders, pigments or additives  
30 conventional in lacquers.

- With regard to the ionic groups, the binders, and the optionally present crosslinking agents, the statements made above in connection with the electrocoating lacquer  
35 compositions (Ic) apply.

An anodic electrocoating lacquer composition is preferably used as the electrocoating lacquer composition (II).

In addition to the base resins and optionally present crosslinking agent, the electrocoating lacquer compositions (II) may contain colouring and/or effect pigments, extenders and/or additives conventional in lacquers. In particular, they contain colouring and/or effect pigments, preferably only colouring absorption pigments, if the second coating layer has an influence on the decorative effect of a laminar structure produced according to the invention. Pigments which may be considered are, for example, conventional inorganic and/or organic coloured pigments and/or effect pigments, such as titanium dioxide, iron oxide pigments, carbon black, phthalocyanine pigments, quinacridone pigments, metal pigments, for example made from titanium, aluminium or copper, interference pigments, such as for example aluminium coated with titanium dioxide, coated mica, graphite effect pigments, lamellar iron oxide or lamellar copper phthalocyanine pigments. Examples of extenders are kaolin, talcum or silicon dioxide.

The pigments may be dispersed to yield pigment pastes, for example using known paste resins. Especially in the case of anodic electrocoating lacquer compositions (II), it is possible to use those pigment pastes as are used in the sprayable aqueous base lacquers familiar to the person skilled in the art for the production of two-layer lacquer coatings of the base lacquer/clear lacquer type.

Conventional additives, as are in particular known for electrocoating lacquer compositions, are possible additives. Examples of these are wetting agents, neutralising agents, levelling agents, catalysts, corrosion inhibitors, anti-foaming agents, solvents, light stabilisers optionally in combination with anti-oxidants.

The second coating layer is preferably applied in a dry film thickness of, for example, 10 to 50  $\mu\text{m}$ , particularly preferably of between 15 and 40  $\mu\text{m}$ , and stoved at elevated temperatures, for example between 130 and 180°C. The second

5 coating layer generally has virtually no electrical conductivity, i.e. in the stoved state it exhibits a specific resistance of in general above  $10^9 \Omega \text{ cm}$ .

The second coating may, but need not, extend over the

10 entire area of the three-dimensional substrate. It is accordingly possible, for example, especially in the second and third embodiments of the process according to the invention, to provide a first coating over the entire area by applying a primer coat of the primer coating composition

15 (Ib) or (Ic) and to provide a second coating of the electrocoating lacquer composition (II) substantially only on the external areas, in particular on the visible surfaces of the substrate to be provided with the laminar structure, i.e. for example not in confined cavities of a

20 vehicle body. Once the second coating layer deposited from the electrocoating lacquer composition (II) has been stoved, the plastic film is applied. This is preferably performed only on external areas, in particular on the visible surfaces of a three-dimensional substrate, for

25 example preferably on the external visible surfaces of an automotive body.

The plastic films applied in the process according to the invention in particular comprise those made from

30 thermoplastics or composite films of two or more plies of one or more different thermoplastics. Examples of suitable thermoplastics are polyolefin plastics, polyamide, polyvinyl chloride. Plastic films made from polyvinyl chloride are preferred. The plastic films generally have a

35 thickness of 20 to 200, preferably of 40 to 100  $\mu\text{m}$  and they determine, either alone or in conjunction with the underlying lacquer layer produced from the coating

composition (II), the decorative effect of the lamellar structures produced according to the invention. The plastic films also contribute towards the protective action of the laminar structures produced according to the invention, for example providing protection from damage by stone impact, in particular when thick plastic films are used.

The plastic films used in the process according to the invention may be non-transparent and cover the substrate completely opaquely, such that the decorative effect of a laminar structure produced according to the invention is determined solely by the plastic film. The plastic film may, for example, be coloured on the surface or consist of a through-coloured plastic. Through-coloured plastics contain an adequate quantity of pigments and plastic films produced therefrom are non-transparent. The plastic films may, however, also additionally be provided with an opaquely pigmented lacquer layer or an opaque print (for example as a pattern or representational image). Pigments which may be considered for through-colouring the plastic or for pigmenting the opaque lacquer layer are conventional inorganic and/or organic colouring and/or effect pigments, such as for example titanium dioxide, carbon black, iron oxide pigments, phthalocyanine pigments, quinacridone pigments, metal pigments, for example made from aluminium, interference pigments, such as for example aluminium coated with titanium dioxide, coated mica or lamellar copper phthalocyanine pigments. Preferably, only absorption pigments are present, i.e. the non-transparent plastic films are preferably plain coloured plastic films. The non-transparent plastic films may have additional lacquer layers and/or be printed (for example with a pattern or image).

Preferably, however, transparent plastic films which determine the decorative effect of the laminar structures produced according to the invention in conjunction with the

coating layer produced from the electrocoating lacquer composition (II) are used in the process according to the invention.

- 5 The transparent plastic films may be colourless or transparently coloured, for example preferably with dyes soluble in the plastic. Colouring may also be provided, optionally also in addition to the use of soluble dyes, by a small, non-opaque quantity of colouring pigments or of
- 10 sufficiently finely divided colouring pigments.

The transparent plastic films may also contain effect pigments.

- 15 The transparent plastic films may also be non-opaquely lacquer coated and/or printed (for example with a pattern or image). Lacquer coatings which may be considered are both clear lacquer layers and single or multi-layer transparent coloured and/or effect lacquer coatings.

- 20 The transparent or non-transparent plastic films used in the process according to the invention may have a structure, for example be embossed, but the plastic films preferably have a smooth, in particular glossy, surface.

- 25 The plastic films are preferably applied in the form of a set of films, i.e. they are preferably in the form of a number of plastic films cut to fit the individual visible surfaces of a three-dimensional substrate. The individual
- 30 plastic films in a set may be of differing designs. In the event that the individual films are of differing designs, substrates may be provided with an overall appearance which is non-uniform with regard to the decorative effect. The individual plastic films of a set preferably have the same
- 35 design.

- The laminar structures are produced according to the invention by applying the plastic films onto the coating layers of the electrocoating lacquer composition (II) which have been deposited and stoved. Application preferably involves an adhesive bond, which may optionally be promoted by suitable measures, for example the action of heat and/or vacuum. Adhesive bonding may be achieved by using a hot-melt adhesive, aqueous dispersion adhesive or a solvent-based adhesive or the plastic films are self-adhesive plastic films provided with a pressure sensitive adhesive on their reverse side, which are applied after removal from a flat backing sheet. Especially where self-adhesive plastic films are used, it is particularly simple, if desired, to renew the outer layer of a laminar structure produced according to the invention by removing the old self-adhesive film and applying a new one. The original design may here be selected or the design may be changed by selecting a different plastic film.
- The plastic films may be adhesively bonded not only in an industrial context, for example by an automotive manufacturer, but also in a trade context, for example in a motor vehicle workshop, or in a do-it-yourself context by the final consumer. Especially in the trade and do-it-yourself context, self-adhesive plastic films are preferably used, particularly preferably a set of self-adhesive plastic films, for example in the case of automotive vehicles with an appropriate number of plastic films cut to fit the individual visible surfaces of the vehicle bodywork.

- The process according to the invention is particularly convenient for providing a surface finish on those three-dimensional substrates, for example automotive bodies or parts thereof, which are of a simple shape, i.e. which have smooth or curved surfaces without numerous corners, edges, beads and angles.

- This invention also relates to the protective and decorative laminar structures applied using the process according to the invention onto the surface of three-dimensional substrates, the laminar structures consisting
- 5 of a primer layer produced from a primer coating composition (Ia) or (Ib) or (Ic), a coating layer of an electrocoating lacquer composition (II) applied thereon and a plastic film adhesively bonded thereon.
- 10 The process according to the invention permits three-dimensional, electrically conductive substrates, in particular automotive bodies, to be provided with a protective and decorative laminar structure. Especially when transparent plastic films are used as the outer layer
- 15 of the laminar structure produced according to the invention, numerous decorative variants may be produced. For example, it is possible to divide the substrates primed using the primer coating composition (Ia) or (Ib) or (Ic) into a number of groups corresponding to the number of
- 20 electrocoating lacquer compositions (II) available with different coloured pigments. Each of the resultant groups of primed substrates may then be provided with a second coating layer of one of the differently pigmented electrocoating lacquer compositions (II). The number of
- 25 electrocoating lacquer compositions (II) used in different shades is preferably limited. Different transparent plastic films may then be applied onto the substrates which have been provided with the second coating layer in one of a limited variety of shades. In this manner, it is possible
- 30 to produce the various possible combinations of transparent plastic films with the available number of differently coloured precoated substrates.
- The process according to the invention is logistically
- 35 straightforward, it may be performed with a limited number of different coating compositions, spray application of coating compositions and thus the consequent formation of



the waste typical of lacquer spraying are avoided. The process according to the invention permits the surface of three-dimensional substrates, in particular motor vehicles, to be provided with a protective coating and decorative finish. Consistent colouring or effects may thus be applied even to assemblies of different substrate components.

Patent Claims

1. A process for the production of a three-dimensional substrate provided with a protective and decorative laminar structure, characterised in that a primer layer of a coating composition (I) which is electrically conductive in the stoved state is applied without spraying onto an electrically conductive substrate and stoved, whereupon a substrate not yet in the desired three-dimensional shape is shaped, whereupon a second coating layer of an electrophoretically depositable coating composition (II) is electrophoretically deposited and stoved and whereupon a plastic film is applied.
2. A process according to claim 1, characterised in that the conductive primer layer is applied by brushing, roller application, dipping or flow coating.
3. A process according to claim 1 or 2, characterised in that the conductive primer layer is applied using the coil coating process by autophoretic deposition or electrophoretic deposition.
4. A process according to any one of claims 1, 2 or 3, characterised in that the conductive primer layer is applied onto the entire surface of the substrate.
5. A process according to any one of claims 1 to 4, characterised in that, once the conductive primer layer has been applied and stoved, the substrate is shaped three-dimensionally and optionally previously stamped or cut.
6. A process according to any one of claims 1 to 5, characterised in that the substrate is in the form of

components which are combined into an assembly before application of the second coating layer.

7. A process according to any one of claims 1 to 5,  
5 characterised in that the conductive primer layer is applied onto both sides of a substrate in the form of a sheet metal coil using the coil coating process and stoved, sheet metal components are stamped out from the coil and shaped and then provided with the second  
10 coating layer by electrophoretic deposition.
8. A process according to any one of claims 1 to 7,  
characterised in that it is used for coating motor  
vehicles or the components thereof.
- 15 9. A decorative laminar structure obtained using the process of any one of claims 1 to 8.
10. A three-dimensional substrate provided with a  
20 protective and decorative laminar structure obtained according to any one of claims 1 to 8.

**Abstract:**

Process for the production of a three-dimensional substrate provided with a protective and decorative laminar structure, in which a primer layer of a coating composition (I) which is electrically conductive in the stoved state is applied without spraying onto an electrically conductive substrate and stoved, whereupon a substrate not yet in the desired three-dimensional shape is shaped, whereupon a second coating layer of an electrophoretically depositable coating composition (II) is electrophoretically deposited and stoved and whereupon a plastic film is applied.

As a below named inventor, I hereby declare that my residence, post office address and citizenship are as stated below next to my name, and I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the INVENTION ENTITLED Process for the application of a protective and laminar structure

the specification of which (CHECK applicable BOX(ES))  
X ☐ is attached hereto.  
BOX(ES) ☐ was filed on \_\_\_\_\_ as U.S. Application No. 0 / \_\_\_\_\_  
☐ was filed as PCT International Application No. PCT/EP98/04614 on 23.07.1998  
and (if U.S. or PCT application amended) was amended on 22.05.1999

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose all information known to me to be material to patentability as defined in 37 C.F.R. 1.56. I hereby claim foreign priority benefits under 35 U.S.C. 119/365 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate filed by me or my assignee disclosing the subject matter claimed in this application and having a filing date (1) before that of the application on which priority is claimed, or (2) if no priority claimed, before the filing date of this application:

PRIOR FOREIGN APPLICATION(S)		Date first Laid- open or Published	Date Patented or Granted	Priority Claimed Yes No
Number	Country	Day/MONTH/Year Filed		
197 33 312.5	DE	01.08.1997	04.02.1999	X

I hereby claim domestic priority benefit under 35 U.S.C. 119/120/365 of the indicated United States applications listed below and PCT international applications listed above or below and, if this is a continuation-in-part (CIP) application, insofar as the subject matter disclosed and claimed in this application is in addition to that disclosed in such prior applications, I acknowledge the duty to disclose all information known to me to be material to patentability as defined in 37 C.F.R. 1.56 which became available between the filing date of each such prior application and the national or PCT international filing date of this application:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

And I hereby appoint Cushman Darby & Cushman Intellectual Property Group of Pillsbury Madison & Suto LLP, 1100 New York Avenue, N.W., Ninth Floor, East Tower, Washington, D.C. 20005-3918, telephone number (202) 861-3000 (to whom all communications are to be directed), and the below-named persons (of the same address) individually and collectively my attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith and with the resulting patent, and I hereby authorize them to delete names/numbers below of persons no longer with their firm and to act and rely on instructions from and communicate directly with the person/assignee/attorney/firm/organization who/which first sends/sent this case to them and by whom/which I hereby declare that I have consented after full disclosure to be represented unless/until I instruct the above Firm and/or a below attorney in writing to the contrary.

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3. INVENTOR'S SIGNATURE: \_\_\_\_\_ Date \_\_\_\_\_

Inventor's Name (typed) \_\_\_\_\_

Residence (City) First \_\_\_\_\_ Middle Initial \_\_\_\_\_ Family Name \_\_\_\_\_ Country of Citizenship

Post Office Address (Include Zip Code) \_\_\_\_\_

(FOR ADDITIONAL INVENTORS, check box ☐ and attach sheet (CDC-116.2) for same information for each re signature, name, date, citizenship, residence and address.)